



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 5 : A61K 37/36, C07K 7/10, 13/00 C07K 3/20	A1	(11) International Publication Number: WO 92/08480 (43) International Publication Date: 29 May 1992 (29.05.92)
(21) International Application Number: PCT/US91/08606		(81) Designated States: AT (European patent), AU, BE (European patent), CA, CH (European patent), DE (European patent), DK (European patent), ES (European patent), FR (European patent), GB (European patent), GR (European patent), IT (European patent), JP, LU (European patent), NL (European patent), SE (European patent).
(22) International Filing Date: 18 November 1991 (18.11.91)		
(30) Priority data: 614,306 16 November 1990 (16.11.90) US		
(71) Applicant: CELTRIX LABORATORIES, INC. [US/US]; 2500 Faber Place, Palo Alto, CA 94303 (US).		Published <i>With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>
(72) Inventors: OWAGA, Yasushi ; 310 Farallon Avenue, Pacifica, CA 94044 (US). SCHMIDT, David ; 3721 Floral Drive, Santa Cruz, CA 95062 (US). DASCH, James ; 837 Seminole Way, Redwood City, CA 94062 (US).		
(74) Agent: CIOTTI, Thomas; Morrison & Foerster, 545 Middlefield Road, Suite 200, Menlo Park, CA 94025 (US).		

(54) Title: A β -TYPE TRANSFORMING GROWTH FACTOR

(57) Abstract

A new heterodimeric form of TGF- β is described. This 25 KD molecule is active in an in vitro assay of inhibition of epithelial cell growth. The protein may be isolated from bone. When reduced, the protein elutes in two peaks by RP-HPLC. In immunoblots, the reduced protein from the earlier eluting peak reacts predominately with antibodies directed against TGF- β 3, while reduced protein from the later eluting peak reacts predominately with antibodies directed against TGF- β 2. The N-terminal amino acid sequence and immunoreactivity of the native protein are consistent with a heterodimer of TGF- β 2 and TGF- β 3.

BEST AVAILABLE COPY

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AT	Austria	ES	Spain	MC	Madagascar
AU	Australia	FI	Finland	ML	Mali
BB	Barbados	FR	France	MN	Mongolia
BE	Belgium	GA	Gabon	MR	Mauritania
BF	Burkina Faso	GB	United Kingdom	MW	Malawi
BG	Bulgaria	GN	Guinea	NL	Netherlands
BJ	Benin	GR	Greece	NO	Norway
BR	Brazil	HU	Hungary	PL	Poland
CA	Canada	IT	Italy	RO	Romania
CF	Central African Republic	JP	Japan	SD	Sudan
CG	Congo	KP	Democratic People's Republic of Korea	SE	Sweden
CH	Switzerland	KR	Republic of Korea	SN	Senegal
CI	Côte d'Ivoire	LI	Liechtenstein	SU+	Soviet Union
CM	Cameroon	LK	Sri Lanka	TD	Chad
CS	Czechoslovakia	LU	Luxembourg	TG	Togo
DE*	Germany	MC	Monaco	US	United States of America
DK	Denmark				

+ Any designation of "SU" has effect in the Russian Federation. It is not yet known whether any such designation has effect in other States of the former Soviet Union.

5

A B-type Transforming Growth FactorTechnical field

10 The present invention relates to protein chemistry. More particularly, it relates to the discovery and isolation of a new form of transforming growth factor β .

Background

15 PCT WO 84/001106, filed 23 Sept. 1983, describes transforming growth factor β_1 (TGF- β_1) and its use for the promotion of cell proliferation and tissue repair, wound healing, and treatment of traumata.

20 U.S. 4,848,063 describes two cartilage inducing factors, CIF-A and CIF-B, found in mammalian bone that (1) are cofactors for inducing cartilage formation *in vivo*; (2) promote connective tissue deposition *in vivo* in the absence of any added activating agent or cofactor, and (3) are active in the anchorage-independent cell growth assay used to characterize TGF- β (this assay is sometimes called the TGF- β assay herein and is described in Methods for Preparation of Media, Supplements, and Substrate for Serum-free Animal Cell Culture (1984) pp. 181-194, Alan R. Liss, Inc.).

30 U.S. 4,806,523, filed 6 March 1986, discloses that CIF-A and CIF-B both possess anti-inflammatory activity and are inhibitors of mitogen stimulated T cell proliferation and B cell activation. It also reports that CIF is localized in centers of hematopoiesis and lymphopoiesis and that CIF may, therefore, be useful for

35

-2-

treating indications associated with malfunction or dysfunction of hematopoiesis or lymphopoiesis. CIF-A has since been shown to be identical to TGF- β 1. CIF-B has since been recognized as a new form of β -type
5 transforming growth factor and is now called TGF- β 2.

U.S. 4,886,747 discloses a third β -type transforming growth factor called TGF- β 3.

TGF- β 1, TGF- β 2, and TGF- β 3 are all composed of two identical polypeptide chains linked by disulfide
10 bonds, i.e., they are homodimers. The heterodimer of TGF- β 1 and TGF- β 2, called TGF- β 1.2, has been identified and its uses demonstrated. PCT WO 88/05788, filed 29 January 1988, discloses a heterodimer of TGF- β 1 and TGF- β 2. PCT WO 90/00900, filed 20 July 1989, discloses
15 treatment of inflammatory disorders with homodimeric TGF- β 1 and 2, and the heterodimer TGF- β 1.2.

Disclosure of the Invention

The invention provides a previously unknown
20 form of TGF- β that is found in bone and methods for obtaining the same in substantially pure form from bone or from recombinant expression in vitro. This TGF- β , denoted TGF- β 2.3, is a heterodimer of TGF- β 2 and TGF- β 3, and is active in an in vitro assay of inhibition of
25 epithelial cell growth.

Accordingly, one aspect of the invention is substantially pure TGF- β 2.3. In another aspect, a chondrogenic/osteogenic effective amount of TGF- β 2.3 with a substantially nonimmunogenic carrier is formulated as a
30 chondrogenic/osteogenic implant composition. In a further aspect an effective amount of TGF- β 2.3 with a pharmaceutically acceptable carrier is formulated as a composition for promoting proliferation of normal cells.

- 3 -

Another aspect of the invention is a process for preparing TGF- β 2.3 from bone, comprising the steps of pooling side fractions from peaks of column chromatography, subjecting those fractions to reverse phase HPLC and recovering those fraction which migrate more slowly than TGF- β 2 by SDS-PAGE, subjecting those slower migrating fractions to FPLC and recovering those that migrate during a pH 4.6 to 6.7 gradient, subjecting the pH 4.6 to 6.7 eluant to reverse phase HPLC or gel electrophoresis, and recovering substantially pure TGF- β 2.3.

Another aspect of the invention is a method of inducing cartilage and/or bone formation at a predetermined site with an implant of TGF- β 2.3.

Another aspect of the invention is a method for treating a patient for osteoporosis with a therapeutically effective amount of a parenteral formulation of TGF- β 2.3 administered parenterally.

Another aspect of the invention is a method for treating a patient for inflammation with an anti-inflammatory effective amount of TGF- β 2.3. A further aspect of the invention is a method for preventing or reducing local inflammation to a solid implant made of a permeable material by dispersing an anti-inflammatory effective amount of TGF- β 2.3 in the material.

An additional aspect of the invention is a method for treating a patient for an indication associated with dysfunction or malfunction of hematopoiesis or lymphopoiesis with an effective amount of TGF- β 2.3.

Another aspect of the invention is a method for inhibiting the growth of tumor cells in a mammal by administering to that mammal an oncostatically effective amount of TGF- β 2.3.

-4-

Another aspect of the invention is a method for producing TGF- β 2.3 by joining the DNA sequence encoding the N-terminal signal sequence and proregion of TGF- β 2 or TGF- β 3 to the DNA sequence encoding the mature sequence of TGF- β 3 or TGF- β 2 to generate a chimeric construct, introducing the chimeric construct in an expression vector into a host cell, introducing a TGF- β 2 or TGF- β 3 precursor gene in an expression vector into the host cell, wherein the TGF- β 2 or TGF- β 3 precursor gene has an N-terminal signal sequence and proregion substantially corresponding to N-terminal signal sequence region in the chimeric construct, and recovering TGF- β 2.3 from the host cell.

A further aspect of the invention is a method for preventing severe cardiac injury resulting from reperfusion of ischemic myocardium comprising administering an effective amount of TGF- β 2.3 to a patient, prior to or after the onset of ischemia.

Another aspect of the invention is a method for the treatment of septic shock in an animal comprising administering an effective amount of TGF- β 2.3 to the animal.

A further aspect of the invention is a method for protecting hematopoietic stem cells in a patient from the myelotoxicity of chemotherapeutic drugs comprising administering an effective amount of TGF- β 2.3 to the patient prior to exposure to the chemotherapeutic drugs.

A further aspect of the invention is a method for protecting hematopoietic stem cells in a patient from the myelotoxicity of radiation therapy comprising administering an effective amount of TGF- β 2.3 to the patient prior to exposure to the radiation therapy.

A further aspect of the invention is a method for diagnosing a disorder involving the production of

-5-

TGF- β 2, TGF- β 3, or TGF- β 2.3, using as a diagnostic reagent TGF- β 2.3 protein, or monoclonal antibodies or polyclonal antibodies directed against TGF- β 2.3.

Another aspect of the invention is a method for
5 treating acute and chronic disease states that result from overproduction of TGF- β 2, 3, or 2.3, by administering a therapeutically effective amount of a monoclonal antibody reactive with TGF- β 2.3, or an antigen-binding fragment of a monoclonal antibody
10 reactive with TGF- β 2.3.

Another aspect of the invention is a method for
treating tumor cells that produce TGF- β 2, 3, or 2.3, by
administering a therapeutically effective amount of a
monoclonal antibody reactive with TGF- β 2.3 to suppress
15 the immunosuppressive effects of TGF- β .

Another aspect of the invention is a method for
treating metastatic cancers by administering a
therapeutically effective amount of a monoclonal antibody
reactive with TGF- β 2.3 to mark tumor cells for
20 destruction by complement or by immune cells dedicated to
tumor removal

These and other embodiments of the present
invention will readily occur to those of ordinary skill
in view of the disclosure herein.

25

Brief Description of the Figures

Figure 1 is the elution profile of TGF- β 2 peak
fractions prepared by cation exchange chromatography,
30 applied to a C18 RP-HPLC column and eluted with a linear
acetonitrile gradient in 0.1% trifluoroacetic acid (TFA).

Figure 2 is the elution profile on a Mono-S
FPLC column in a pH 4.6 to 6.7 and pH 6.7 to 9.0 gradient
of a TGF- β 2 peak fraction, prepared as in Figure 1, the

-6-

pH 4.6 to 6.7 gradient fraction having a slightly slower mobility in SDS-PAGE than other TGF- β 2 peak fractions.

Figure 3 is the elution profile of a fraction containing predominately 25 KD protein obtained from a pH 5 4.6 to pH 6.7 shift on a Mono-S FPLC column, chromatographed on a C18 reverse phase HPLC column in a linear acetonitrile gradient. The protein eluted as one major peak with a minor peak overlapping on the backside.

Figure 4 is a comparison of the elution 10 profiles of TGF- β 2.3 with TGF- β 1 and TGF- β 2 from a C18 reverse phase HPLC column in an acetonitrile gradient in 0.1% TFA.

Figure 5 is the elution profile of reduced TGF- β 2.3 as two peaks from a C18 reverse phase HPLC column in 15 an acetonitrile gradient in 0.1% TFA, the later eluting peak overlapping with the peak of reduced TGF- β 2.

Detailed Description

The practice of the present invention will 20 employ, unless otherwise indicated, conventional techniques of synthetic organic chemistry, protein chemistry, molecular biology, microbiology, and recombinant DNA technology, which are within the skill of the art. Such techniques are explained fully in the literature. See, e.g., Scopes, R.K., Protein Purification Principles and Practices, 2d ed. (Springer-Verlag, 1987), Methods in Enzymology (S. Colowick and N. Kaplan, eds., Academic Press, Inc.), Sambrook et al., Molecular Cloning: A Laboratory Manual, 2d ed., Cold 25 Spring Harbor Press, Cold Spring Harbor, NY, 1989, Handbook of Experimental Immunology, Vols. I-IV (D.M. Weir and C.C. Blackwell, eds, 1986, Blackwell Scientific Publications); House, Modern Synthetic Reactions, 2nd ed., Benjamin/Cummings, Menlo Park, Cal., 1972.

- 7 -

All patents, patent applications, and publications mentioned herein, whether supra or infra, are hereby incorporated by reference in their entirety.

5 A. Definitions

In defining the present invention, the following terms will be employed, and are intended to be defined as indicated below.

As used herein the term "treat" is intended to mean prophylaxis or attenuation of an existing condition. Accordingly, in the case of inflammation, the invention method may be used to prevent inflammation or alleviate existing inflammation.

As used herein the term "inflammation" is intended to encompass both acute responses (i.e. a response in which the inflammatory processes are active) and chronic responses (i.e. a response marked by slow progress and formation of new connective tissue).

Chronic and acute inflammation may be distinguished by the cell types involved. Acute inflammation often involves polymorphonuclear neutrophils; whereas chronic inflammation is normally characterized by a lymphohistiocytic and/or granulomatous response.

Examples of specific types of inflammation are diffuse inflammation, focal inflammation, croupous inflammation, interstitial inflammation, obliterative inflammation, reactive inflammation, specific inflammation, toxic inflammation, and traumatic inflammation.

As used herein, the term "septic shock" refers to the sequence of events triggered by bacteremia during which cell wall substances (endotoxin in Gram-negative organisms and peptidoglycan/teichoic acid complex in Gram-positive organisms) activate the complement, kinin,

- 8 -

and ACTH/endorphin systems. This series of metabolic events ultimately progresses to a state of shock.

As used herein, a protein is substantially pure when that protein has been purified to the extent that it is essentially free of other molecules with which it is associated in nature. In this regard, the term "substantially pure" intends a composition containing less than about 30% by weight contaminating protein, preferably less than about 10% contaminating protein, and most preferably less than about 5% by weight contaminating protein. The term "substantially pure" is used relative to proteins with which the TGF- β 2.3 is associated in nature and is not intended to exclude compositions in which the TGF- β 2.3 is admixed with nonproteinaceous pharmaceutical carriers or vehicles or proteinaceous pharmaceutical carriers or vehicles.

As used herein, an amino acid sequence substantially corresponding to TGF- β 2 will have at least 80% sequence homology with the amino acid sequence of TGF- β 2. Similarly, an amino acid sequence substantially corresponding to TGF- β 3 will have at least 80% sequence homology with the amino acid sequence of TGF- β 3.

As used herein, a DNA sequence substantially corresponding to TGF- β 2 will have at least 80% sequence homology with the DNA sequence of TGF- β 2. Similarly, a DNA sequence substantially corresponding to TGF- β 3 will have at least 80% sequence homology with the DNA sequence of TGF- β 3.

As used herein, an "expression vector" refers to a plasmid, bacteriophage, virus, or other molecule into which a gene of interest may be cloned, such that the appropriate signals for expression of that gene are present on that vector. Expression vectors may require regulation of expression by factors provided in trans.

-9-

As used herein, the terms "monoclonal antibody" and "Mab" refer to an immunoglobulin composition having a substantially homogeneous population of antibodies, each of which binds to the same antigenic determinant. Unless otherwise indicated, the term is not intended to be limited to antibodies of any particular mammalian species or isotype, or to antibodies prepared in any given manner. The term is intended to include whole antibody molecules as well as antigen-binding fragments (e.g., Fab', F(ab')₂).

10 B. General Methods

The present invention relates to TGF- β 2.3, a novel β -type transforming growth factor. TGF- β s isolated to date from natural sources are polypeptide 15 dimers of approximately 25 to 26 KD molecular weight as determined by SDS-PAGE. TGF- β 2.3 is a heterodimer of TGF- β 2 and TGF- β 3.

A procedure for isolating TGF- β s from bovine bone is described in U.S. Patent No. 4,843,063, which is 20 incorporated herein by reference in its entirety. It involves extracting demineralized bone (DMB) with an extractant (e.g. \geq 4M guanidine hydrochloride, 8M urea) that solubilizes nonfibrous proteins, gel filtering the extract to obtain a <30 KD fraction on 25 carboxymethylcellulose (CMC) at pH 4.5-5.5, preferably 4.8, eluting the CMC-absorbed fraction with an NaCl gradient, and purifying the proteins from the portion eluting at about 150-250 mM NaCl by RP-HPLC or gel electrophoresis.

30 TGF- β s exhibit activity in the TGF- β assay described in Methods for Preparation of Media, Supplements, and Substrate for Serum-free Animal Cell Culture (1984) pp 181-194, Alan R. Liss, Inc. That assay determines ability to induce anchorage-independent growth

-10-

in non-neoplastic normal rat kidney fibroblasts by measuring the formation of cell colonies in soft agar. Procedures for obtaining TGF- β s from platelets, placenta, and kidney tissues are described in International patent

5 publication WO 84/01106 and EPA publication number 0128849. Briefly, this involves extracting the source material with acid-ethanol, sizing the extract by gel filtration, and isolating the TGF- β from the filtrate by high performance liquid chromatography (HPLC).

10 TGF- β s isolated to date are nonspecies specific as regards TGF- β activity. It is believed, therefore, that these polypeptides have been highly conserved among animal species (i.e., a given polypeptide from different mammalian species has an amino acid sequence that varies, 15 if at all, in one or more amino acid residue deletions, additives, or substitutions that do not affect the non-species specific activity of the molecule adversely) and have cross-species functionality. For example, murine TGF- β 1 has been shown to differ from human TGF- β 1 by only 20 one amino acid (Derynck et al., J. Biol. Chem. 261:4377, 1986), while murine TGF- β 2 differs from human TGF- β 2 by only three amino acids (Miller et al., Mol. Endo. 3:1108, 1989). Bovine TGF- β 1 and TGF- β 2 have completely 25 identical amino acid sequences to human TGF- β 1 and TGF- β 2, respectively (Ogawa and Seyedin, Meth. Enz. Vol 198, in press). Human TGF- β 3 differs from chicken TGF- β 3 by only one amino acid residue, but is identical to murine TGF- β 3 (Denhez et al., Growth Factors 3:139, 1990). Accordingly, the TGF- β s may be derived from cells or 30 tissue of diverse animal origin or may be obtained by recombinant DNA technology. Correlatively, TGF- β from one vertebrate species may be used to treat another vertebrate species. The most common therapeutic usages of TGF- β will be in the treatment of humans, domestic

-11-

animals such as cattle, sheep, and pigs, and sports or pet animals such as dogs, cats, and horses.

The TGF- β 2.3 of the invention may be useful by itself or in combination with cofactors for inducing cartilage/bone formation for repairing, replacing, or augmenting cartilage/bone tissue in animals including humans. Chondrogenically/osteogenically effective amounts of the proteins will normally be formulated with pharmacologically and physiologically acceptable fluid or solid carriers for implantation.

The TGF- β 2.3 of the present invention may also be useful as a diagnostic reagent for detecting cancers, neoplasms, and other disorders involving the production of TGF- β 2, TGF- β 3, or TGF- β 2.3. Purified TGF- β 2.3 may be used as the reagent itself, or alternatively, monoclonal or polyclonal antibodies directed against TGF- β 2.3 may be used as the diagnostic reagent in such diagnostic assays.

The TGF- β 2.3 of the invention may also be used in the same manner as other TGF- β s to promote (provoke and sustain) non-species specific cellular proliferation. Clinical applications of the cell proliferation activity of these compositions include topical administration for burn or wound healing or tissue repair. In such uses the proteins and activating agent will be formulated in amounts sufficient to induce soft tissue cell proliferation with pharmaceutically acceptable carriers that are added for the particular mode of administration. Topical dosage forms will typically be formulated as sprays, gels, ointments, or salves. Implants will be formulated as injectables. Systemic dosage forms may be formulated for enteral administration (i.e., liquids, pills, tablets) or for parenteral injection. The dosages used in such applications cannot be specified because of

-12-

the nature of all proliferation activity and the variability in wounds and other trauma.

The TGF- β 2.3 may also be useful for treating bone deficiencies, such as osteoporosis and

5 osteopetrosis, systemically. For such treatment the TGF- β 2.3 will be formulated in therapeutically effective amounts with injectable carriers and administered parenterally to the patient. Doses will typically be in the range of 0.001 μ g/kg to 10 mg/kg.

10 TGF- β s may be used as oncostats in treating any type of cellular neoplasm, including, without limitation, carcinomas, myelomas, melanomas, and lymphomas.

Particularly preferred targets are breast, lung, colon, and ovarian carcinomas. The TGF- β s may be administered

15 locally or systemically, depending on the nature and degree of the neoplasm being treated. For local administration an oncostatically effective amount of TGF- β 2.3 or mixtures formulated thereof with a pharmaceutically acceptable carrier as an injectable for 20 parenteral administration, or as a solid or semisolid implant which may or may not be of a sustained or a controlled release form.

Alternatively, the oncostats could be delivered to solid tumors in particular, including inoperable

25 tumors using current catheter technology for localized delivery via the arterial supply to the tumor. In this situation the oncostat could be mixed with a vasoocclusive agent, such as injectable collagen, which would provide a means to reduce perfusion of the tumor and at the same 30 time provide for the localized delivery of the oncostatic agent. Clips may also be used to occlude venous drainage, and thus maintain high doses of TGF- β 2.3 in the tumor mass.

-13-

For systemic administration, oncostatically effective amounts of TGF- β 2.3 will be formulated with conventional carriers used for water soluble proteins (e.g. physiological saline, sugar solutions and the like) for injection into circulation. Alternatively, they may be formulated as a sustained release formulation that releases the TGF- β 2.3 to circulation over a prolonged time period. Specific targeting of the factor for tumor cells in systemic applications may be accomplished by conjugation of the TGF- β 2.3 to an antibody directed against tumor specific cell surface antigen(s). Enhanced tumor cell cytotoxicity may be accomplished by covalently radiolabelling the TGF- β 2.3 with ^{131}I , a cytotoxic agent. The TGF- β s are readily iodinated and retain full biological activity. Monoclonal antibody preparations with specificity for particular tumor types, such as breast and ovarian tumors, are well known in the art. Other oncostats of chemotherapeutic drugs may be included in the formulation if desired.

The term "oncostatically effective" is intended to indicate a dose that effects a significant (>50%) inhibition of tumor cell proliferation. In *in vitro* assays, 50% inhibition is generally observed at TGF- β concentrations of the order of 0.2 $\mu\text{g}/\text{ml}$ or lower and saturation is achieved at 10 $\mu\text{g}/\text{ml}$ or lower. Inhibition may be monitored *in vivo* by monitoring the patient's tumor burden. The amount of TGF- β 2.3 which is oncostatically effective in a given treatment will depend upon the patient, the type and degree of cancer being treated and the mode of administration. In general, the amounts administered to adult humans will be in the range of about 0.001 $\mu\text{g}/\text{kg}$ to 10 mg/kg. Corresponding systemic administration will involve the higher segment of the

-14-

range (0.01 $\mu\text{g}/\text{kg}$ to 10 mg/kg) due to the clearance or other in situ inactivation of the polypeptide.

The TGF- β 2.3 of the instant invention may be useful in the treatment of both local and systemic inflammation. When used as a local anti-inflammatory agent the TGF- β 2.3 will usually be formulated in effective amounts with pharmaceutically acceptable carriers in weight ratios to carrier in the range of 1:1000 to 1:20000. If tissue deposition at the site is not desired, the level of TGF- β 2.3 to carrier may be lowered to below that (e.g. at weight ratios below 1:6000 in the case of collagen carrier which promotes tissue deposition. In addition to being formulated as an injectable, the TGF- β 2.3 may be incorporated (dispersed) into solid permeable implants such as collagenous soft and hard tissue implants, prostheses, sponges, wound dressings, and sutures to modulate local inflammatory responses to solid bodies. Since such implants are made from permeable materials the TGF- β 2.3 can diffuse from the implant and exert its anti-inflammatory properties.

When used to treat inflammation at internal sites locally, the TGF- β 2.3 may be injected, inhaled, placed surgically, or otherwise administered locally, depending on the particular formulation, and the site where inflammation control is desired.

For systemic administration TGF- β 2.3 may be formulated with conventional carriers used with water soluble proteins for injection into circulation. Alternatively, it may be formulated as a sustained release implant formulation if the indication being treated so requires.

The amount of TGF- β 2.3 administered to treat inflammation will depend upon the patient, the inflammatory condition being treated, and the mode of

-15-

administration. In general, amounts administered to adult humans will be in the range of about 0.001 µg/kg to 10 mg/kg. When the TGF- β 2.3 is administered locally, amounts in the lower portion of the range will normally be used, typically 0.1 to 10 µg. Correspondingly, systemic administration will typically involve amounts in the 10-1000 µg range.

TGF- β 2.3 may be particularly effective in the treatment of inflammation involving the respiratory system. In this application, the TGF- β 2.3 may be administered by inhalation with a suitable aerosol. In this form, these factors would be useful for the treatment of diffuse interstitial diseases of the lung such as asbestosis, silicosis, or coal-miner's pneumoconiosis; the treatment of immunological diseases that involve the respiratory tract such as rheumatoid arthritis, lupus erythematosus, or Goodpasture's syndrome; and the treatment of granulomatosis and eosinophilic granulomatosis.

These anti-inflammatory peptides may be combined with carriers in the form of a salve, ointment, or other topical formulation and thereby be useful in the control of dermal inflammation by topical application. Such formulations would be particularly useful in the local treatment of psoriasis vulgaris, contact dermatitis, dermal ulcers, and acute or chronic eczematous dermatitis.

TGF- β 2.3 may be either used alone or combined with a slow release carrier and injected into or around joints, bone, or muscle for control of inflammation associated with various diseases. Some examples include myositis (viral, bacterial, parasitic, fungal, or autoimmune processes); myasthenia gravis; osteomyelitis; osteoarthritis and rheumatoid arthritis.

-16-

Since TGF- β molecules have been shown to be stable at low pH and resistant to enzyme digestion, these factors may be delivered systemically by ingestion. These properties make these factors particularly useful
5 for the treatment of gastric and duodenal ulcers, granulomatous gastritis, esophagitis (numerous causes); enteritis (numerous causes); and colitis (numerous causes).

TGF- β s have also been shown to be effective for
10 the treatment of septic shock (International Publication Number WO 90/000903, filed 21 July, 1989). TGF- β 2.3 may be administered prophylactically or therapeutically, i.e., before, simultaneous with, or after an infection has set in. TGF- β 2.3 may be used to treat patients who
15 are at-risk with respect to bacterial infection, or who suffer from septicemia. Patients who are at risk include those receiving immunosuppressive therapy and those suffering from severe thermal burns or other serious injury, cystic fibrosis, renal failure, or cancer, or who
20 are undergoing extensive surgical procedures or organ transplantation.

TGF- β 2.3 may be administered to these patients by any suitable technique, including systemic or local, as discussed above. Similarly, TGF- β 2.3
25 compositions and doses may be formulated as discussed for above applications, taking into consideration the requirements of the individual patient, the cause of the septic shock, the method of administration, and other factors known to practitioners. Doses will typically be
30 in the range of 0.001 μ g/kg to 10 mg/kg.

TGF- β 2.3 may be used to treat a patient for an indication associated with a dysfunction or malfunction of hematopoiesis or lymphopoiesis. TGF- β 2.3 may be administered to these patients by any suitable

-17-

technique, including systemic or local, as discussed above. Similarly, TGF- β 2.3 compositions and doses may be formulated as discussed for above applications, taking into consideration the requirements of the individual patient, the method of administration, and other factors known to practitioners. Doses will typically be in the range of 0.001 $\mu\text{g}/\text{kg}$ to 10 mg/kg.

TGF- β may also be used to protect hematopoietic stems cells from the myelotoxicity of chemotherapeutic drugs, such as cyclophosphamide and mephalan, or radiation therapy. In such applications a therapeutically effective amount of TGF- β 2.3 will be administered prior, usually 3-72 hours, to the administration of the chemotherapeutic drug or radiation therapy. The mode of administration is preferably interfemoral arterial, intraperitoneal, or subcutaneous, and is preferably by injection. TGF- β 2.3 compositions and doses may be formulated as discussed for above applications, taking into consideration the requirements of the individual patient, the nature of the drug or radiation therapy used, the method of administration of the composition, and other factors known to practitioners. Doses will typically be in the range of 0.001 $\mu\text{g}/\text{kg}$ to 10 mg/kg.

TGF- β may also be used in the prevention of severe cardiac injury resulting from reperfusion of ischemic myocardium (Lefer et al., Science 249:61, 1990). TGF- β 2.3 may be administered, preferably intravenously, prior to or after the onset of ischemia. TGF- β 2.3 compositions and doses may be formulated as discussed for above applications, taking into consideration the requirements of the individual patient and other factors known to practitioners. Doses will typically be in the range of 0.001 $\mu\text{g}/\text{kg}$ to 10 mg/kg.

Recombinant Expression of TGF- β 2.3

The proteins of the invention may be expressed in vitro, or in vivo in either prokaryotic or eukaryotic systems. Prokaryotes are most frequently represented by various strains of E. coli. However, other microbial strains may also be used, such as bacilli (for example Bacillus subtilis), various species of Pseudomonas, and other bacterial strains. In such prokaryotic systems, plasmid vectors which contain replication sites and control sequences derived from a species compatible with the host are used. For example, E. coli is typically transformed using derivatives of pBR322, a plasmid derived from an E. coli species by Bolivar et al., Gene 2:95, 1977. Commonly used prokaryotic control sequences, which are defined herein to include promoters for transcription initiation, optionally with an operator, along with ribosome binding site sequences, include such commonly used promoters as the β -lactamase (penicillinase) and lactose (lac) promoter systems (Chang et al., Nature 198:1056, 1977) and the tryptophan (trp) promoter system (Goeddel et al., Nuc. Acids Res. 8:4057, 1980) and the lambda-derived P_L promoter and N-gene ribosome binding site (Shimatake et al., Nature 292:128, 1981). However, any available promoter system compatible with prokaryotes can be used.

The expression systems useful in eukaryotic systems of the invention comprise promoters derived from appropriate eukaryotic genes. A class of promoters useful in yeast, for example, include promoters for synthesis of glycolytic enzymes, including those for 3-phosphoglycerate kinase (Hitzeman et al., J. Biol. Chem. 255:2073, 1980). Other promoters include those from the enolase gene (M.J. Holland et al., J. Biol. 35

-19-

Chem. 256:1385, 1981) or the Leu2 gene obtained from YEpl3 (J. Broach et al., Gene 8:121, 1978).

Suitable mammalian promoters include the early and late promoters from SV40 (Fiers et al., Nature 273:113, 1978) or other viral promoters such as those derived from polyoma, adenovirus II, bovine papilloma virus, or avian sarcoma viruses. Suitable viral and mammalian enhancers are cited above. In the event plant cells are used as an expression system, the nopaline synthesis promoter is appropriate (Depicker et al., J. Mol. Appl. Gen. 1:561, 1982). Expression in insect cell culture may conveniently be achieved using a baculovirus vector.

Construction of suitable vectors containing the desired coding and control sequences employs standard ligation and restriction techniques which are well understood in the art. Isolated plasmids, DNA sequences, or synthesized oligonucleotides are cleaved, tailored and religated in the form desired.

Site specific DNA cleavage is performed by treatment with a suitable restriction enzyme (or enzymes) under conditions which are generally understood in the art, generally following the manufacturer's directions. See, e.g., New England Biolabs, Product Catalog. In general, about 1 µg of plasmid or DNA sequence is cleaved by one unit of enzyme in about 20 µl of buffer solution; in the examples herein, typically, an excess of restriction enzyme is used to insure complete digestion of the DNA substrate. Incubation times of about 1 hr to 2 hr at about 37°C are workable, although variations can be tolerated. After each incubation, protein may be removed by extraction with phenol/chloroform, and may be followed by diethyl ether extraction, and the nucleic acid recovered from aqueous fractions by ethanol precipi-

-20-

tation followed by separation over a Sephadex® G-50 spin column. If desired, size separation of the cleaved fragments may be performed by polyacrylamide gel or agarose gel electrophoresis using standard techniques. A general description of size separations is found in Meth. Enzymol. 65:499-560, 1980.

Single-strand "ends" of restriction cleaved fragments may be removed or converted to a double-strand form ("blunt-ended") by treating with the large fragment of E. coli DNA polymerase I (Klenow) in the presence of the four deoxyribonucleotide triphosphates (dNTPs) using incubation times of about 15 to 25 min at 20 to 25°C in 50 mM Tris, pH 7.6, 50 mM NaCl, 6 mM MgCl₂, 6 mM DTT and 5-10 µM dNTPs. The Klenow fragment adds deoxyribonucleotides at 5' "sticky ends" but hydrolyzes protruding 3' single strands, even though the four dNTPs are present. If desired, selective repair can be performed by supplying only 1-3 of the dNTPs, within the limitations dictated by the nature of the sticky ends.

After treatment with Klenow fragment, the mixture is extracted with phenol/chloroform and ethanol precipitated followed by chromatography on a Sephadex G-50 column. Treatment under appropriate conditions with S1 nuclease results in hydrolysis of any single-stranded portion.

Synthetic oligonucleotides are prepared by the triester method of Matteucci et al. (J. Am. Chem. Soc. 103:3185, 1981) or using commercially available automated oligonucleotide synthesizers. Labeling of single strands prior to annealing is achieved using an excess, e.g., approximately 10 units of polynucleotide kinase to 0.1 nmole substrate in the presence of 50 mM Tris, pH 7.6, 10 mM MgCl₂, 5 mM dithiothreitol, 1-2 mM ATP, 1.7 pmoles ³²P-ATP (2.9 mCi/mmmole), 0.1 mM spermidine, and 0.1 mM EDTA.

-21-

Ligations are performed in 15-30 μ l volumes under the following standard conditions and temperatures: 20 mM Tris-HCl, pH 7.5, 10 mM MgCl₂, 10 mM DTT, 33 μ g/ml BSA (bovine serum albumin), 10 mM-50 mM NaCl, and either 5 40 μ M ATP, 0.01-0.02 (Weiss) units T4 DNA ligase at 0°C (for "sticky end" ligation) or 1 mM ATP, 0.3-0.6 (Weiss) units T4 DNA ligase at 14°C (for "blunt end" ligation). Intermolecular "sticky end" ligations are usually 10 performed at 33-100 μ g/ml total DNA concentrations (5-100 nM total end concentration). Intermolecular blunt end ligations (usually employing a 10-30 fold molar excess of linkers) are performed at 1 μ M total ends concentration.

Correct ligations for plasmid construction may be confirmed by first transforming E. coli strain MM294 15 obtained from E. coli Genetic Stock Center, CGSC #6135, or other suitable host with the ligation mixture. Successful transformants are selected by ampicillin, tetracycline or other antibiotic resistance or using other markers depending on the plasmid construction, as 20 is understood in the art. Plasmids from the transformants are then prepared according to the method of D.B. Clewell et al, Proc. Natl. Acad. Sci. U.S.A. 62:1159, 1969, optionally following chloramphenicol amplification (D.B. Clewell, J. Bacteriol. 110:667, 1972). 25 The isolated DNA is analyzed by restriction and/or sequenced by the dideoxy method of F. Sanger et al., Proc. Natl. Acad. Sci. U.S.A. 74:5463, 1977, as further described by Messing et al., Nucleic Acids Res. 9:309, 1981, or by the method of Maxam et al., Meth. Enzymol. 30 65:499, 1980.

In vector construction employing "vector fragments", the vector fragment is commonly treated with bacterial alkaline phosphatase (BAP) in order to remove the 5' phosphate and prevent religation of the vector.

-22-

BAP digestions are conducted at pH 8 in approximately 150 mM Tris, in the presence of Na⁺ and Mg²⁺ using about 1 unit of BAP per µg of vector at 60°C for about 1 hr. In order to recover the nucleic acid fragments, the 5 preparation is extracted with phenol/chloroform and ethanol precipitated and desalted by application to a Sephadex G-50 column. Alternatively, religation can be prevented in vectors which have been double-digested by additional restriction enzyme digestion of the unwanted 10 fragments.

For portions of vectors derived from cDNA or genomic DNA which require sequence modifications, site-specific primer directed mutagenesis may be used. This is conducted using a synthetic oligonucleotide primer 15 complementary to a single-stranded phage DNA to be mutagenized except for limited mismatching, representing the desired mutation. Briefly, the synthetic oligonucleotide is used as a primer to direct synthesis of a strand complementary to the phage, and the resulting 20 double-stranded DNA is transformed into a phage-supporting host bacterium. Cultures of the transformed bacteria are plated in top agar, permitting plaque formation from single cells which harbor the phage.

Theoretically, 50% of the new plaques will 25 contain the phage having the mutated form as a single strand; 50% will have the original sequence. The resulting plaques are hybridized with kinased synthetic primer under allele-specific conditions. In general, one may vary the temperature, ionic strength, and concentration 30 of chaotropic agent(s) in the hybridization solution to obtain conditions under which substantially no probes will hybridize in the absence of an "exact match." For hybridization of probes to bound DNA, the

-23-

empirical formula for calculating optimum temperature under standard conditions (0.9 M NaCl) is

$$T(^{\circ}\text{C}) = 4(N_G + N_C) + 2(N_A + N_T) - 5^{\circ}\text{C},$$

5

where N_G , N_C , N_A , and N_T are the numbers of G, C, A, and T bases in the probe (J. Meinkoth et al., Anal. Biochem. 138:267, 1984). Plaques which hybridize with the probe are then picked, cultured, and the DNA recovered.

10

When a vector is used that requires transformation of the host cell, transformation of the host cell with a recombinant construct is done using standard techniques appropriate to such cells. The calcium treatment employing calcium chloride, as

15

described by Cohen, Proc. Natl. Acad. Sci. U.S.A.

69:2110, 1972, or the RbCl method described in Maniatis et al., Molecular Cloning: A Laboratory Manual (1982)

Cold Spring Harbor Press, p. 254, is used for prokaryotes or other cells which contain substantial cell wall

20

barriers. Infection with Agrobacterium tumefaciens (Shaw et al., Gene 23:315, 1983) is used for certain plant cells. For mammalian cells without cell walls, the calcium phosphate precipitation method of Graham and van der Eb, Virology 52:546, 1978 is preferred.

25

Transformations into yeast are carried out according to the method of Van Solingen et al., J. Bac. 130:946, 1977, and C.L. Hsiao et al., Proc. Natl. Acad. Sci. U.S.A.

76:3829, 1979. Alternatively, one may use a liposomal transfection system. For example, one may use a

30

synthetic lipid such as N-[1-(2,3-dioleyloxy)-propyl]-N,N,N-trimethylammonium chloride, commercially available under the name Lipofectin (BRL, Gaithersburg, MD), as described by P.L. Felgner et al., Proc. Natl. Acad. Sci. U.S.A. 84:7413, 1987.

35

-24-

The cDNA or genomic libraries resulting from any of the transformation methods described above are screened using the colony hybridization procedure. Each microtiter plate is replicated onto duplicate nitro-cellulose filter papers (S & S type BA-85) and colonies are allowed to grow at 37°C for 14-16 hr on L agar containing 50 µg/ml ampicillin. The colonies are lysed and DNA fixed to the filter by sequential treatment for 5 min with 500 mM NaOH, 1.5 M NaCl, and are washed twice for 5 min each time with 5X standard saline citrate (SSC). Filters are air dried and baked at 80°C for 2 hr. The duplicate filters are prehybridized at 42°C for 6-8 hr with 10 ml per filter of DNA hybridization buffer (5X SSC, pH 7.0, 5X Denhardt's solution (polyvinyl-pyrrolidone, plus Ficoll and bovine serum albumin; 1X = 0.02% of each), 50 mM sodium phosphate buffer at pH 7.0, 0.2% SDS, 20 µg/ml poly-U, and 50 µg/ml denatured salmon sperm DNA).

The samples are hybridized with kinased probe under conditions which depend on the stringency desired. Typical moderately stringent conditions employ a temperature of 42°C for 24-36 hr with 1-5 ml/filter of DNA hybridization buffer containing probe. For higher stringencies, high temperatures and shorter times are employed. The filters are washed four times for 30 min each time at 37°C with 2X SSC, 0.2% SDS and 50 mM sodium phosphate buffer at pH 7, then washed twice with 2X SSC and 0.2% SDS, air dried, and are autoradiographed at -70°C for 2 to 3 days.

Probes for detection of the desired recombinant construct may be prepared as follows. Nucleic acid binding partners are prepared by means known to those of ordinary skill in the art, for example by cloning and restriction of appropriate sequences or preferably by

-25-

direct chemical synthesis. For example, one may employ the phosphotriester method described by S.A. Narang et al., Meth. Enzymol., 68:90, 1979, and U.S. Pat. No. 4,356,270, incorporated herein by reference.

5 Alternatively, one may use the phosphodiester method disclosed in E.L. Brown et al., Meth. Enzymol. 68:109, 1979, incorporated herein by reference. Other methods include the diethylphosphoramide method disclosed in Beaucage et al., Tetrahedron Lett., 22:185, 1981, and the 10 solid support method disclosed in U.S. Pat. No. 4,458,066. The binding partners may also be labeled, if desired, by incorporating means detectable by spectroscopic, photochemical, biochemical, immunochemical, or chemical means. For example, the 15 primer may include ³²P, fluorescent dyes, electron-dense reagents, enzymes (as commonly used in ELISAs), biotin, or haptens or proteins for which antisera or monoclonal antibodies are available.

Antibodies to the proteins of the invention, 20 both polyclonal and monoclonal, may be prepared by conventional methods. In general, the protein is first used to immunize a suitable animal, preferably a mouse, rat, rabbit or goat. Rabbits and goats are preferred for the preparation of polyclonal sera due to the volume of 25 serum obtainable, and the availability of labeled anti-rabbit and anti-goat antibodies. Immunization is generally performed by mixing or emulsifying the protein in saline, preferably in an adjuvant such as Freund's complete adjuvant, and injecting the mixture or emulsion 30 parenterally (generally subcutaneously or intramuscularly). A dose of 50-200 µg/injection is typically sufficient. Immunization is generally boosted 2-6 weeks later with one or more injections of the protein in saline, preferably using Freund's incomplete adjuvant.

-26-

One may alternatively generate antibodies by in vitro immunization using methods known in the art, which for the purposes of this invention is considered equivalent to in vivo immunization.

5 Polyclonal antisera is obtained by bleeding the immunized animal into a glass or plastic container, incubating the blood at 25°C for one hour, followed by incubating at 4°C for 2-18 hours. The serum is recovered by centrifugation (e.g., 1,000 x G for 10 minutes).

10 About 20-50 ml per bleed may be obtained from rabbits.

Monoclonal antibodies are prepared using the method of Kohler and Milstein, Nature 256:495, 1975, or a modification thereof. Typically, a mouse or rat is immunized as described above. However, rather than 15 bleeding the animal to extract serum, the spleen (and optionally several large lymph nodes) is removed and dissociated into single cells. If desired, the spleen cells may be screened (after removal of nonspecifically adherent cells) by applying a cell suspension to a plate or well coated with the protein antigen. B cells expressing membrane-bound immunoglobulin specific for the antigen bind to the plate, and are not rinsed away with the rest of the suspension. Resulting B cells, or all dissociated spleen cells, are then induced to fuse with 20 myeloma cells to form hybridomas, and are cultured in a selective medium (e.g., hypoxanthine, aminopterin, thymidine medium, "HAT"). The resulting hybridomas are plated by limiting dilution, and are assayed for the production of antibodies which bind specifically to the 25 immunizing antigen (and which do not bind to unrelated antigens). The selected monoclonal antibody (MAb)-secreting hybridomas are then cultured either in vitro (e.g., in tissue culture bottles or hollow fiber reactors), or in vivo (as ascites in mice).

-27-

If desired, the antibodies (whether polyclonal or monoclonal) may be labeled using conventional techniques. Suitable labels include fluorophores, chromophores, radioactive atoms (particularly ^{32}P and ^{125}I), electron-dense reagents, enzymes, and ligands having specific binding partners. Enzymes are typically detected by their activity. For example, horseradish peroxidase is usually detected by its ability to convert 3,3',5,5'-tetramethylbenzidine (TMB) to a blue pigment, quantifiable with a spectrophotometer. "Specific binding partner" refers to a protein capable of binding a ligand molecule with high specificity, as for example in the case of an antigen and a monoclonal antibody specific therefor. Other specific binding partners include biotin and avidin or streptavidin, IgG and protein A, and the numerous receptor-ligand couples known in the art. It should be understood that the above description is not meant to categorize the various labels into distinct classes, as the same label may serve in several different modes. For example, ^{125}I may serve as a radioactive label or as an electron-dense reagent. Horseradish peroxidase (HRP) may serve as enzyme or as antigen for a MAb. Further, one may combine various labels for desired effect. For example, MAbs and avidin also require labels in the practice of this invention: thus, one might label a MAb with biotin, and detect its presence with avidin labeled with ^{125}I , or with an anti-biotin MAb labeled with HRP. Other permutations and possibilities will be readily apparent to those of ordinary skill in the art, and are considered as equivalents within the scope of the instant invention.

Antibodies reactive with TGF- β neutralize the biological activity of TGF- β by preventing the antigen from binding to its cell surface receptors. The intact

-28-

antibody, antigen binding fragments (e.g., Fab', F(ab')₂) may be useful in these applications. In addition, administration of anti-TGF- β 2.3 antibodies would form immune complexes (antigen-antibody) complexes that will 5 increase the rate at which the antigen TGF- β is cleared from the systemic circulation or from the tissue site where the antigen is produced.

Fibrotic diseases and tumor cells may be treated by administering a therapeutically effective 10 amount of anti-TGF- β 2.3 antibodies to affect the inhibition of fibrosis formation or regression of tumor cells. The method and frequency of administration, the dose range, and the duration of antibody therapy will vary with the severity and nature of the condition, and 15 the general health of the patient.

In a preferred embodiment, the antibodies of the present invention are administered locally to the affected tissue sites by bolus injection or perfusion. The amount of antibody administered may be measured by 20 maintaining the local tissue concentration of TGF- β at about 1-1000 μ g/ml.

Indications where this mode of treatment is particularly useful are for the control of excessive scar tissue formation, due to surgery or trauma, or prevention 25 of the formation of connective tissue adhesions. For the treatment of tumor cells by local administration, the antibodies may be delivered directly into a solid tumor mass through a vascular catheter for deep solid tumors, or through a hypodermic needle for superficial or 30 cutaneous tumors. The antibodies may be locally administered by a single bolus injection that is repeated over several days, or by continuous administration by perfusion. The amount of antibody administered is preferably about 1 μ g up to 1000 μ g/g tumor tissue.

-29-

In another embodiment, the antibodies may be administered systemically by intravenous or peritoneal perfusion, or by bolus injection into the subcutaneous tissue or muscle. The antibody may be delivered in vehicles generally known to those skilled in the art, such as saline, balanced salt solution, isotonic or phosphate buffered saline (pH 7), with or without dextrose.

Indications where this mode of treatment is particularly useful are systemic diseases such as interstitial lung fibrosis, liver cirrhosis, scleroderma, and metastatic cancer.

For both local and systemic administration, antibodies reactive with TGF- β 2.3 may be administered in combination with other antibodies reactive with TGF- β to reduce the amount of bioavailable factor.

Experimental

The following examples are intended to illustrate specific embodiments of the invention. They are not intended to limit the invention in any matter.

A. Isolation of TGF- β 2.3

Bovine bone was processed as described in, for example, U.S. 4,843,063, to purify TGF- β 2. After cation exchange chromatography, fractions containing the peak of TGF- β 2 were identified and pooled.

This pool was applied onto a C18 reverse-phase HPLC column (1 x 25 cm, 5 μ , Vydac, 218TP510) and the bound proteins were eluted with a linear acetonitrile gradient in 0.1% TFA (Figure 1). SDS-PAGE of the fractions revealed that the majority of the fractions contained a major band migrating at 25 KD. However, in a

-30-

fraction (fraction #5) which eluted between the peaks of TGF- β 1 and TGF- β 2, the band migrated slightly slower than in the other fractions. Fraction #5 was applied onto a Mono-S FPLC column (0.5 X 5 cm, Pharmacia, HR5/5), which
5 had been equilibrated into 6 M urea, 50 mM sodium acetate, 10 mM NaCl, 1% isopropanol, pH 4.6. The column was equilibrated into 6 M urea, 50 mM sodium acetate, 10 mM NaCl, 1% isopropanol, pH 6.7, over a 10 minute period at a flow rate of 0.5 ml/min to raise the pH to 6.7. The
10 column was then equilibrated into 6M urea, 20 mM HEPES, 10mM NaCl, 1% isopropanol, pH 6.7. Finally, the column was equilibrated into the buffer of the same composition, but at pH 9.0, over 10 minutes at flow rate of 0.5 ml/min. Some protein eluted during the pH 4.6 to 6.7
15 gradient, but the majority of the TGF- β 2 eluted during the pH 6.7 to 9.0 gradient (Figure 2).

SDS-PAGE analysis of the pH 4.6-6.7 fractions revealed that fraction #4 consisted of predominately 25 KD protein. This #4 fraction containing the 25 KD protein in the pH 4.6-6.7 gradient was chromatographed on a C18 reverse-phase HPLC column (0.46 x 25 cm, Vydac, 5 μ , 218TP54). The protein was eluted with a linear acetonitrile gradient in 0.1% TFA. The protein eluted as one major peak (peak #1) with a minor peak (peak #2)
25 overlapping on the back side of the major peak (Figure 3). SDS-PAGE showed that both major and minor peaks contained one single band at 25 KD under non-reducing conditions and 12 KD to 13 KD under reducing conditions, which are consistent with TGF- β .

30 TGF- β 1 eluted before TGF- β 2 from the C18-reverse phase HPLC columns during acetonitrile gradient in 0.1% TFA. Under identical conditions, recombinant TGF- β 3 has been reported to elute after TGF- β 2 (Graycar et al., Mol Endo. 3: 1977-1986, 1989). The TGF- β 2.3
35

-31-

heterodimer (peak #1) eluted as a single sharp peak slightly before TGF- β 2, but after TGF- β 1, under identical conditions (Figure 4). When the protein was reduced with β -mercaptoethanol prior to HPLC, the protein then eluted
5 in two peaks (Figure 5). The position of the later eluting peak coincided with the position of the TGF- β 2 peak when reduced. Results from immunoblots demonstrated that the later eluting peak contained predominately TGF- β 2, while the earlier eluting peak contained
10 predominately TGF- β 3. The results were consistent with the identity of the 25 KD peak as a TGF- β 2.3 dimer.

B. Cell Culture Assay

Mink lung epithelial cells (Mv1Lu, ATCC CCL 64)
15 were cultured on 96 well tissue culture plates at a concentration of 1×10^3 cells per 50 μ l MEM containing 10% fetal calf serum, penicillin, streptomycin, non-essential amino acids, and L-glutamine. Test samples were diluted in the culture medium and appropriate
20 dilutions made. Samples of the diluted cells (50 μ l) were added to the test wells in triplicate 30 minutes after plating. After incubation at 37° C in a humidified 5% CO₂ - 95% air atmosphere for 4 days, the wells were rinsed with PBS (phosphate buffered saline)
25 and filled with 100 μ l of 0.1 M sodium acetate, pH 5.5, 0.1% Triton X-100, 10 mM p-nitrophenol phosphate to lyse the cells. The growth of the cells was measured by assaying a constitutively expressed enzyme, acid phosphatase. After 2 hr, 10 μ l of 0.1 N NaOH was added
30 to each well. Absorbance was measured on a multichannel plate reader at an absorbance setting of 405 nm and a reference filter setting of 492 nm. Inhibition-stimulation of growth was expressed in per cent decrease

-32-

of the acid phosphatase activity in the treated cells when compared to the activity in untreated cells.

The protein from peaks #1 and #2 from the reverse-phase HPLC inhibited proliferation of mink lung epithelial cells (Mv1Lu, ATCC CCL64) with an ED₅₀ of 0.02 ng/ml and 0.025 ng/ml, respectively. TGF- β 1 and TGF- β 2 have an ED₅₀ of 0.01 - 0.02 ng/ml. The result demonstrates that the TGF- β 2.3 of the present invention has specific biological activity comparable to TGF- β 1 and TGF- β 2.

C. Amino terminus sequencing

The amino terminus of the protein of peak #1 was sequenced to residue 59. The sequence consisted of an equal mixture of TGF- β 2 and TGF- β 3 amino terminal residues (Table I). The TGF- β 1 sequence was absent. Given a mixture of TGF- β 2 with either TGF- β 1 or TGF- β 3, distinction between TGF- β 1 and TGF- β 3 can be made at residues # 9, 10, 11, 13, 19, 26, 33, 40, 45, 51, 52, 54, 57, and 58 within the first 59 residues. At residues # 11, 12, 13, 17, 33, 40, and 50, the signals of the residues that match TGF- β 3 were present at approximately 50% level of the total peak signal (the other 50% is from TGF- β 2). Beyond residue #40, signals became lost in the background and could not be identified definitively. Based on these comparisons of partial sequences of N-termini, molecular weight analyses, and biological activity, TGF- β 2.3 appeared to consist of a polypeptide having substantial homology at the N-terminal amino acid sequence to TGF- β 2 and a polypeptide having substantial homology at the N-terminal amino acid sequence to TGF- β 3, both polypeptides being present in equal proportion.

Bovine TGF- β 2.3 N-Terminal Sequence

Ala-leu-Asp-Ala-Ala-Tyr-Xxx¹-Phe-Arg-Asn
Thr Asn

Val-Gln-Asp-Asn-Xxx¹-Xxx²-Leu-Arg-Pro-Leu
Leu Glu Glu Val

Tyr-Ile-Asp-Phe-Lys-Arg-Asp-Leu-Gly-Xxx²
Arg Gln

Lys-Trp-Ile-His-Glu-Pro-Lys-Gly-Tyr-Asn
Val Tyr

Ala-Asn-Phe-Xxx¹-Ala-Gly-Ala-Xxx¹-Xxx³-Tyr
Ser Pro Pro

Leu-Trp-Ser-Ser-Asp-Thr-Xxx⁵-Xxx⁶-Xxx⁷
Tyr Xxx⁴ Arg Tyr Ala
Leu Leu Gly
Thr Thr Ile
Leu Leu
Pro Pro
Tyr Tyr
Val Val

Xxx¹=Cys

Xxx⁴=Arg

Xxx⁷=Ser

Xxx²=Trp

Xxx⁵=Gln/Thr

Xxx³=Pro

Xxx⁶=His

Table 1

3. Immunological characterization

The TGF- β 2.3 protein of the present invention cross-reacted strongly with anti-TGF- β 3 polyclonal antibody on immunoblots. The antibody does not recognize

5 TGF- β 2. The TGF- β 2.3 of the present invention also cross-reacted strongly with 3C7.14.6 anti-TGF- β 2 monoclonal antibody on immunoblots. The monoclonal antibody does not recognize TGF- β 1. These results demonstrated that the TGF- β 2.3 contained TGF- β 2 and TGF-

10 β 3 epitopes.

Furthermore, the TGF- β 2.3 cross-reacted as well as TGF- β 2 with the 2G1 anti-TGF- β 2 monoclonal antibody in an enzyme-linked immunosorbent assay (ELISA). The monoclonal antibody did not cross-react with TGF- β 1 or 15 TGF- β 3, while the antibody cross-reacted only partially with a mixture of TGF- β 2 and TGF- β 3 homodimers. These results demonstrate that the TGF- β 2.3 of the present invention is a heterodimer and not a mixture of TGF- β 2 and TGF- β 3 homodimers.

20

E. Expression of recombinant constructs to yield the heterodimer

The mature biologically active TGF- β 2.3 heterodimer may be produced by cloning and expressing the full-length nucleotide sequence encoding the TGF- β 2 and the TGF- β 3 precursors or their functional equivalent in a host cell which processes the precursor correctly, so 25 that a mature TGF- β 2.3 heterodimer is produced having a biological activity that is virtually indistinguishable from that of the naturally occurring TGF- β 2.3 heterodimer. Functional equivalents of the full length 30 nucleotide sequence encoding the TGF- β 2 and the TGF- β 3 precursors include any DNA sequence which, when expressed 35

inside an appropriate host cell, is capable of directing the synthesis, processing, and export of mature TGF- β 2.3 heterodimer.

Hybrid precursor sequences encoding, for example, the TGF- β 2 precursor sequence joined in-frame to the TGF- β 3 mature sequence in place of the TGF- β 2 mature sequence may be constructed and cloned into an appropriate host cell. A TGF- β 2 precursor sequence with the TGF- β 2 mature sequence is also cloned into the very same host cell to express both the hybrid and the TGF- β precursor sequences in that host cell. Simultaneous expression of the two constructs, consisting of one TGF- β 2 precursor gene and one hybrid gene consisting of the TGF- β 2 N-terminal signal peptide sequence and pro-region joined to TGF- β 3 mature sequence, results in association of the two gene products and expression and production of the TGF- β 2.3 heterodimer.

Similarly, a TGF- β 3 precursor sequence joined in-frame to the mature TGF- β 2 mature sequence in place of the TGF- β 3 mature sequence, may be constructed and cloned into an appropriate host cell. More generally, cloning and expression of any two hybrid precursor sequences, consisting of mature TGF- β 2 and TGF- β 3 sequences joined to separate, but identical signal peptide and TGF- β pro-region sequences in the same host cell, result in production of the TGF- β 2.3 heterodimer.

In a further method for production of recombinant TGF- β 2.3, TGF- β 2 and TGF- β 3 monomeric peptide chains may be renatured separately to produce folded TGF- β monomers. Subsequently, these TGF- β 2 and TGF- β 3 monomers are bonded together through one or more cysteine disulfide bond pairs to form the TGF- β 2.3 heterodimer. For this purpose, the TGF- β gene encoding the mature TGF- β sequence is cloned into a bacterial expression system,

such as E. coli. TGF- β 2 and TGF- β 3 polypeptide chains are expressed separately in different expression systems. TGF- β 2 and TGF- β 3 polypeptide chains are purified and renatured to fold the peptide chain and to form 5 intrachain disulfide bonds. TGF- β 2 and TGF- β 3 monomers are then joined together through an interchain disulfide bond to form the TGF- β 2.3 heterodimer.

Thus, a new form of a TGF- β -type factor, its
10 uses, and methods for its production, have been disclosed. Although the preferred embodiments of the subject invention have been described in some detail, it is understood that obvious variations can be made without departing from the spirit and the scope of the invention
15 as defined by the appended claims.

20

25

30

35

Claims

1. Substantially pure TGF- β 2.3.

5

2. A chondrogenic/osteogenic material comprising:

(a) a chondrogenically or osteogenically effective amount of TGF- β 2.3, and

10

(b) a substantially nonimmunogenic carrier.

3. A composition for promoting proliferation of normal animal cells comprising:

15

(a) an effective amount of TGF- β 2.3; and

(b) a pharmaceutically acceptable carrier.

4. A composition for promoting connective tissue deposition comprising:

20

an effective amount of TGF- β 2.3; and

a pharmaceutically acceptable carrier.

5. A process for preparing TGF- β 2.3 from bone which comprises:

25

(a) pooling TGF- β 2 peak fractions prepared by cation-exchange column chromatography;

(b) subjecting the pool of step (a) to reverse phase HPLC and recovering a fraction which migrates more slowly than TGF- β 2 by SDS-PAGE;

30

(c) subjecting the slower migrating fraction of step (b) to FPLC recovering proteins which elute during a pH 4.6 to 6.7 gradient;

(d) subjecting the pH 4.6 to 6.7 eluant from step (c) to reverse phase HPLC or nondenaturing gel electrophoresis; and

35

(e) recovering substantially pure TGF- β 2.3 from the reverse phase HPLC or nondenaturing gel electrophoresis.

5 6. A protein with TGF- β like activity, wherein said protein is prepared by the process of claim 5.

10 7. A method of inducing cartilage and/or bone formation at a predetermined site in a living mammal comprising implanting TGF- β 2.3 at said site.

15 8. A method of treating a patient for osteoporosis comprising administering a therapeutically effective amount of a parenteral formulation of TGF- β 2.3 to the patient parenterally.

20 9. A method of treating a patient for inflammation comprising administering an anti-inflammatory effective amount of TGF- β 2.3 to a patient.

10. The method of claim 9, wherein the inflammation is acute.

25 11. The method of claim 9 wherein the inflammation is chronic.

12. The method of claim 9 wherein the TGF- β 2.3 is administered systemically.

30 13. The method of claim 9 wherein the TGF- β 2.3 is administered locally.

14. The method of claim 9 wherein the inflammation involves the respiratory system of the patient.

5 15. The method of claim 14 wherein the protein is administered to the patient by inhalation.

10 16. The method of claim 9 wherein the inflammation is dermal inflammation.

17. The method of claim 16 wherein the TGF- β 2.3 is administered topically.

15 18. The method of claim 9 wherein the inflammation is associated with an autoimmune disease.

19. The method of claim 9 wherein the inflammation is in the gastrointestinal tract.

20 20. The method of claim 9 wherein the TGF- β 2.3 is administered in combination with a pharmaceutically acceptable carrier.

25 21. The method of claim 9 wherein the carrier is a collagenous material and the weight ratio of protein to carrier is in the range of 1:1000 to 1: 20000.

30 22. The method of claim 9 wherein the TGF- β 2.3 is substantially free of any activation agent or cofactor.

35 23. A method of preventing or reducing the local inflammatory response to a solid implant made of a permeable material comprising dispersing an anti-

inflammatory effective amount of TGF- β 2.3 in the material.

24. The method of claim 23, wherein said amount is below that which promotes tissue deposition.

25. The method of claim 23, wherein the TGF- β 2.3 is substantially free of any activating agent or cofactor.

10

26. The method of claim 23, wherein the material is collagenous.

15

27. A method for treating a patient for an indication associated with dysfunction or malfunction of hematopoiesis or lymphopoiesis comprising administering an effective amount of TGF- β 2.3 to a patient.

20

28. A method for inhibiting growth of tumor cells in a mammal comprising administering an oncostatically effective amount of TGF- β 2.3 to the mammal.

25

29. The method of claim 28, wherein the TGF- β 2.3 is of the same mammalian species as the mammal.

30. The method of claim 28, wherein the mammal is a human.

30

31. The method of claim 28, wherein the tumor cells are carcinoma cells, adenocarcinoma cells, melanoma cells, or lymphoma cells.

35

32. The method of claim 28, wherein the carcinoma cells are breast, lung, colon, or ovarian carcinoma.

5 33. A method of promoting proliferation of normal cells in an animal comprising administering the composition of claim 3 to an animal.

10 34. A method of promoting proliferation of soft tissue cells in an animal comprising administering the composition of claim 3 to the animal.

15 35. A method of promoting connective tissue deposition at a predetermined site in a patient comprising placing the composition of claim 4 at said site.

36. A method for producing TGF- β 2.3, comprising
20 joining the DNA sequence encoding the N-terminal signal sequence and proregion of TGF- β 2 or TGF- β 3 to the DNA sequence encoding the mature sequence of TGF- β 3 or TGF- β 2 to generate a chimeric construct; introducing the chimeric construct in an expression vector into a host cell;
25 introducing a TGF- β 2 or TGF- β 3 precursor gene in an expression vector into the host cell, wherein the TGF- β 2 or TGF- β 3 precursor gene has an N-terminal signal sequence and proregion substantially corresponding to N-terminal signal sequence region in the chimeric construct;
30 recovering TGF- β 2.3 from said host cell.

37. The method of claim 36, wherein the chimeric construct and the TGF- β 2 or TGF- β 3 precursor molecule are carried on the same expression vector.

5 38. The method of claim 36, wherein the chimeric construct and the TGF- β 2 or TGF- β 3 precursor molecule are carried on the same expression vector.

10 39. The method of claim 36, wherein the expression vector is inducible.

40. The method of claim 36, wherein the host cell is eukaryotic.

15 41. The chimeric construct of claim 36.

20 42. A method for diagnosing a disorder involving the production of TGF- β 2 or TGF- β 3, wherein a diagnostic reagent is selected from the group comprising purified TGF- β 2.3, monoclonal antibodies directed against TGF- β 2.3, or polyclonal antibodies directed against TGF- β 2.3.

25 43. A method for the treatment of septic shock in an animal comprising administering an effective amount of TGF- β 2.3 to the animal.

30 44. A method for protecting hematopoietic stem cells in a patient from the myelotoxicity of chemotherapeutic drugs comprising administering an effective amount of TGF- β 2.3 to the patient prior to exposure to said chemotherapeutic drugs.

45. A method for protecting hematopoietic stem cells in a patient from the myelotoxicity of radiation therapy comprising administering an effective amount of TGF- β 2.3 to the patient prior to exposure to said radiation therapy.

5 46. A method for preventing severe cardiac injury resulting from reperfusion of ischemic myocardium comprising administering an effective amount of TGF- β 2.3 to the patient, prior to or after the onset of ischemia.

10 47. A method for preventing acute or chronic fibrosis, comprising administering a therapeutically effective amount of a monoclonal antibody reactive with TGF- β 2.3

15 48. The method of claim 47, wherein the antibody reactive with TGF- β 2.3 binds an epitope on TGF- β , thereby blocking the binding of TGF- β to TGF- β cellular receptors.

20 49. A method for neutralizing the inhibitor effects of TGF- β , which method comprises administering a therapeutically effective amount of an antibody reactive with TGF- β 2.3.

25 50. A method for inducing tumor regression in a mammal having tumor cells which produce TGF- β , which method comprises administering an amount of a monoclonal antibody reactive with TGF- β 2.3 sufficient to reduce the immunosuppressive effects of TGF- β produced by said tumor cells.

1/5

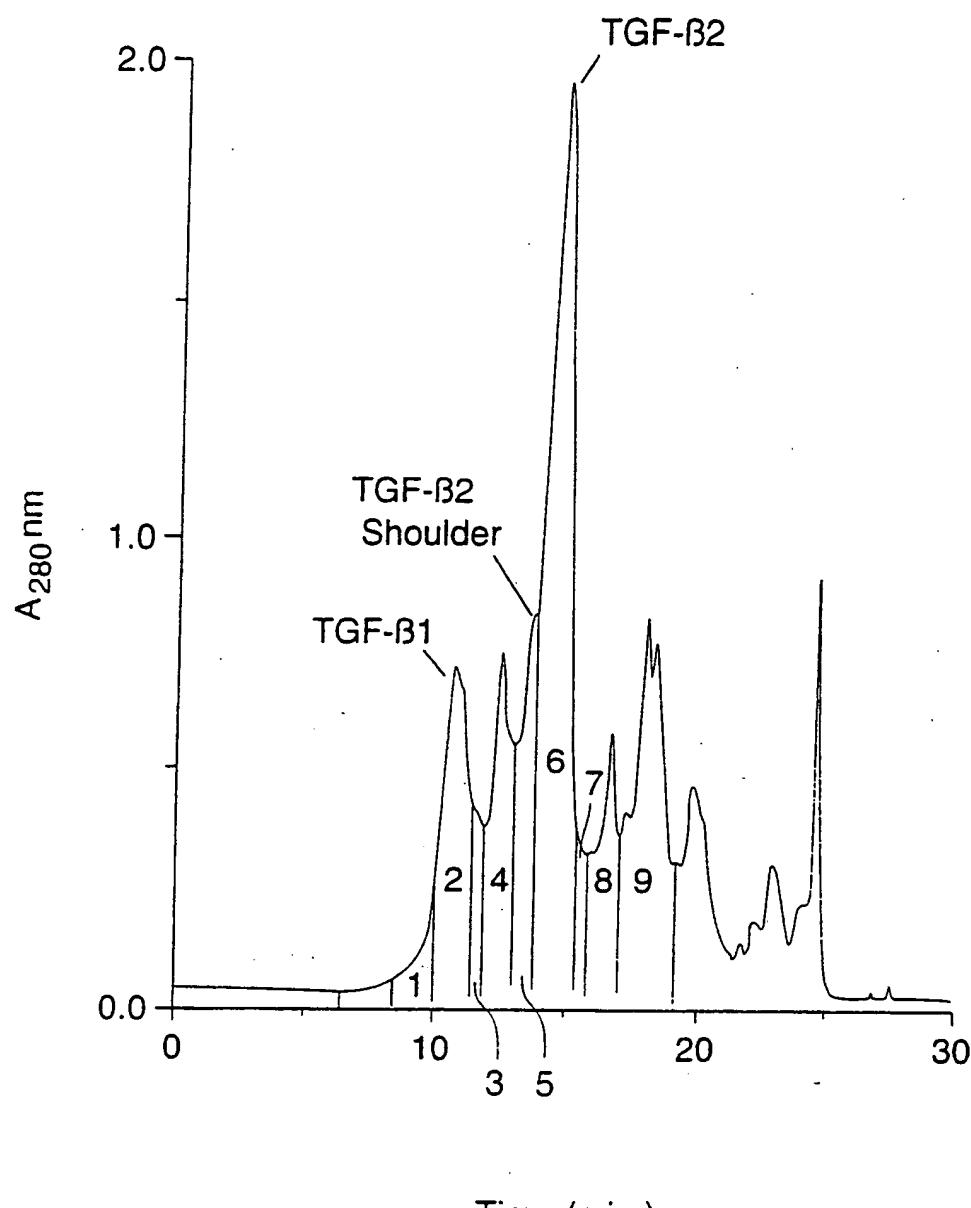
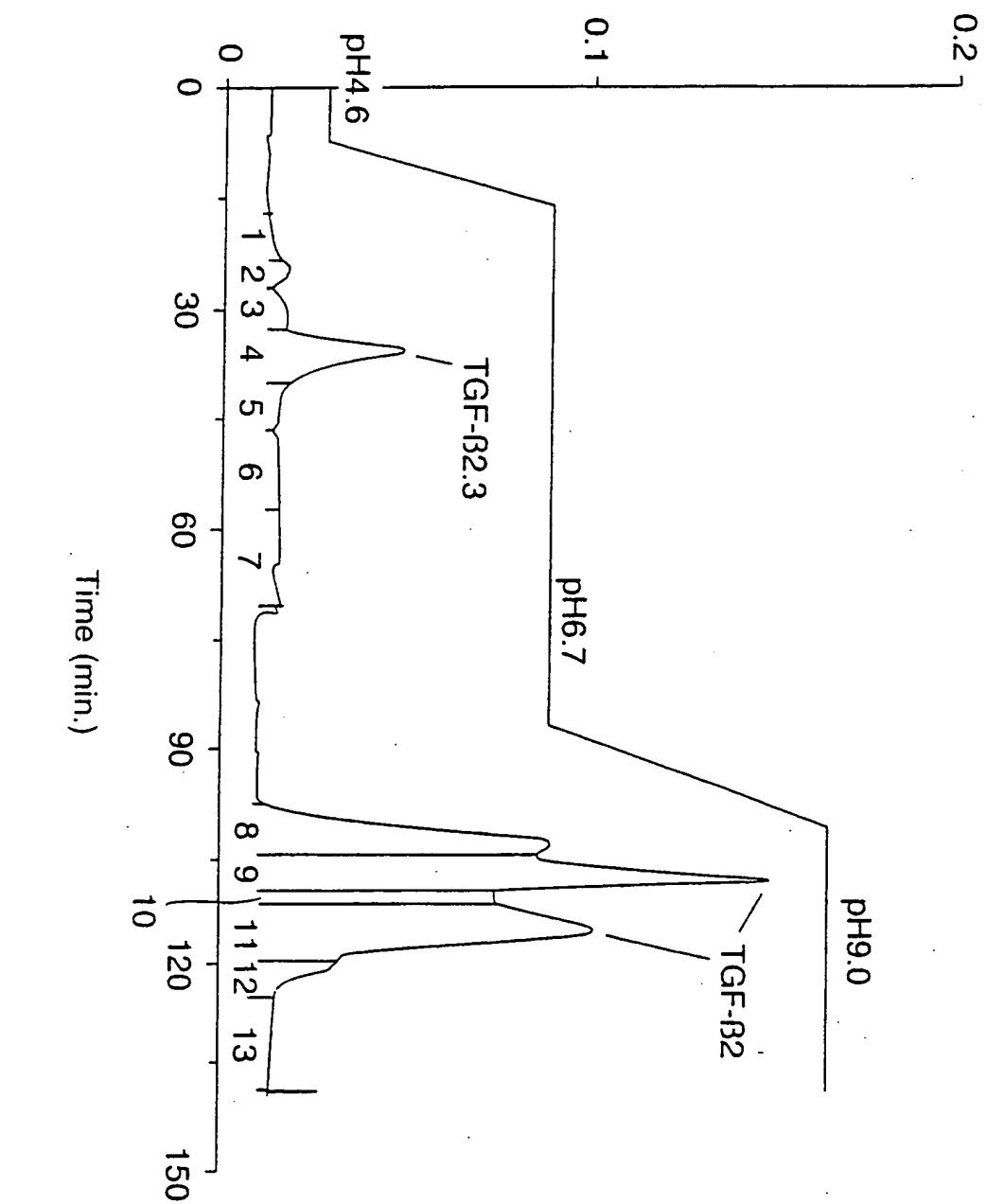


Figure 1

2/5

 $A_{280\text{ nm}}$ 

SUBSTITUTE SHEET

Figure 2

3/5

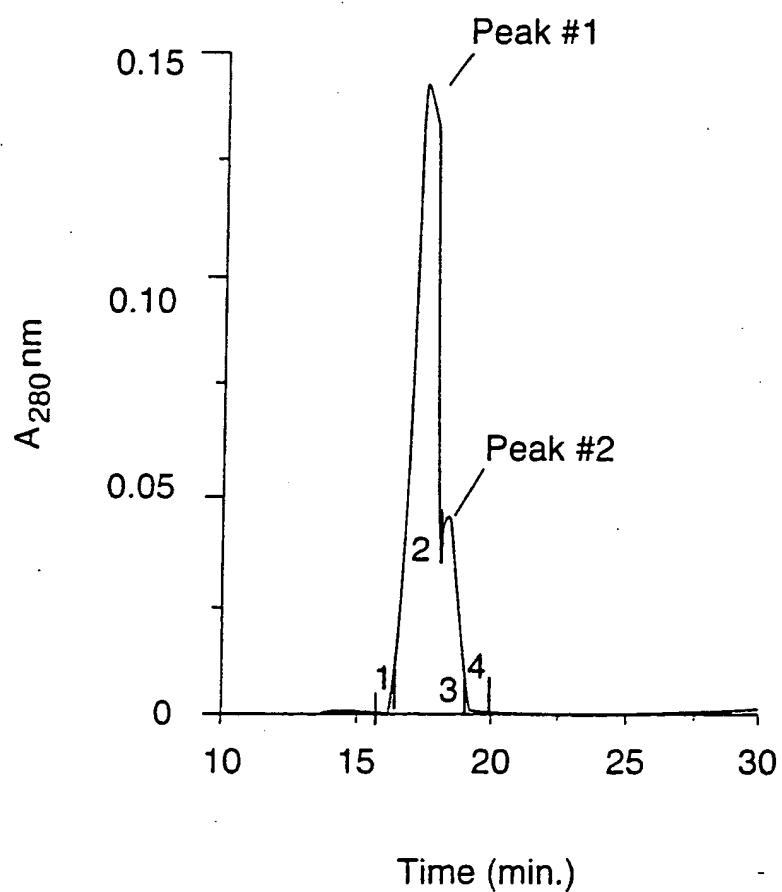


Figure 3

4/5

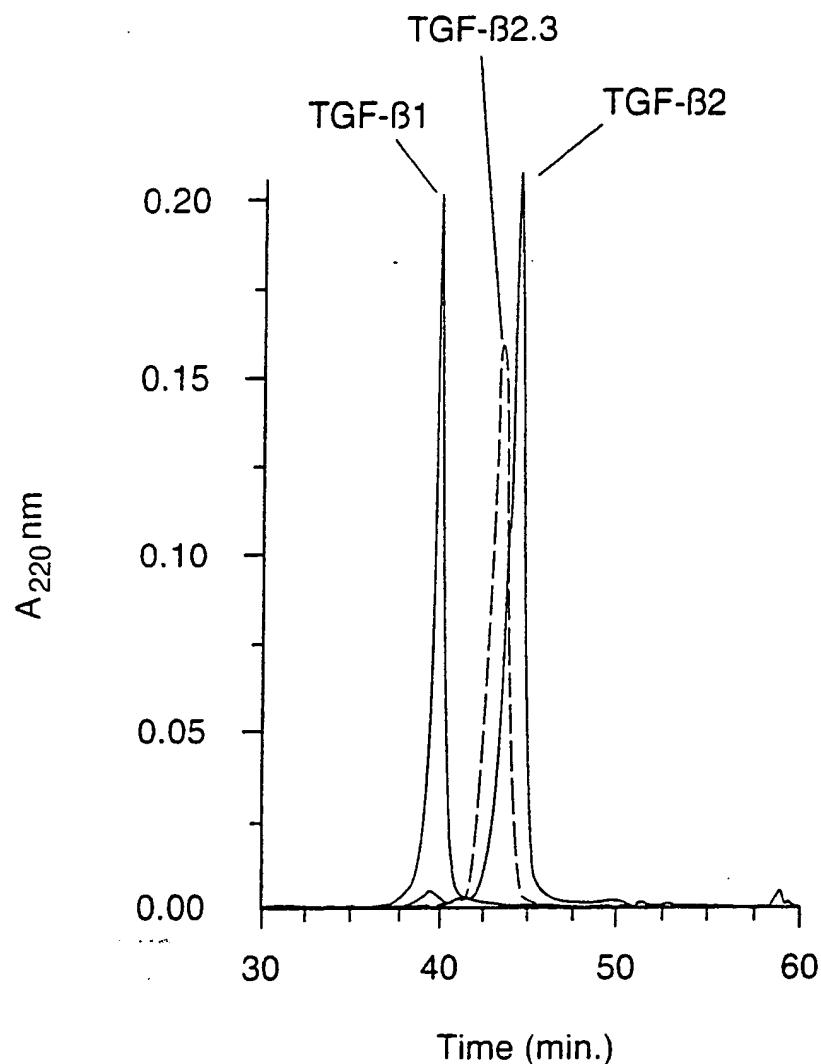


Figure 4

5/5

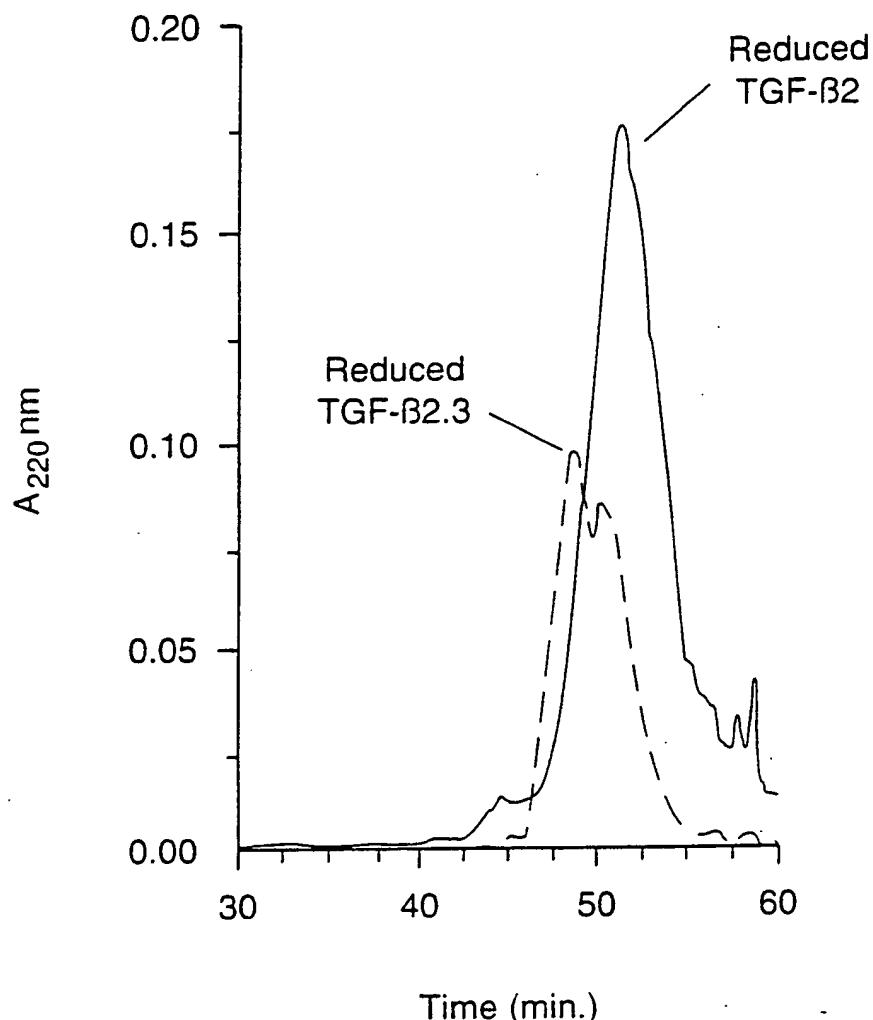


Figure 5

REVISED
VERSION*

PCT

WORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 5 : A61K 37/36, 35/32, C07K 7/10, 13/00 C07K 3/20, 3/28, 15/06, C12P 21/00	A1	(11) International Publication Number: WO 92/08480 (43) International Publication Date: 29 May 1992 (29.05.92)
(21) International Application Number: PCT/US91/08606		(81) Designated States: AT (European patent), AU, BE (European patent), CA, CH (European patent), DE (European patent), DK (European patent), ES (European patent), FR (European patent), GB (European patent), GR (European patent), IT (European patent), JP, LU (European patent), NL (European patent), SE (European patent).
(22) International Filing Date: 18 November 1991 (18.11.91)		
(30) Priority data: 614.306 16 November 1990 (16.11.90) US		
(71) Applicant: CELTRIX LABORATORIES, INC. [US/US]; 2500 Faber Place, Palo Alto, CA 94303 (US).		Published <i>With a revised version of the international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>
(72) Inventors: OWAGA, Yasushi ; 310 Farallon Avenue, Pacifica, CA 94044 (US). SCHMIDT, David ; 3721 Floral Drive, Santa Cruz, CA 95062 (US). DASCH, James ; 837 Seminole Way, Redwood City, CA 94062 (US).		
(74) Agent: CIOTTI, Thomas; Morrison & Foerster, 545 Middlefield Road, Suite 200, Menlo Park, CA 94025 (US).		(88) Date of publication of the revised version of the international search report: 6 August 1992 (06.08.92)

(54) Title: A β -TYPE TRANSFORMING GROWTH FACTOR

(57) Abstract

A new heterodimeric form of TGF- β is described. This 25 KD molecule is active in an in vitro assay of inhibition of epithelial cell growth. The protein may be isolated from bone. When reduced, the protein elutes in two peaks by RP-HPLC. In immunoblots, the reduced protein from the earlier eluting peak reacts predominately with antibodies directed against TGF- β 3, while reduced protein from the later eluting peak reacts predominately with antibodies directed against TGF- β 2. The N-terminal amino acid sequence and immunoreactivity of the native protein are consistent with a heterodimer of TGF- β 2 and TGF- β 3.

FOR THE PURPOSES OF INFORMATION ONLY

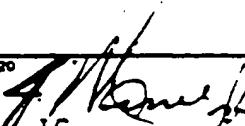
Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AT	Austria	FI	Finnland	MI	Mali
AU	Australia	FR	France	MN	Mongolia
BB	Bahamas	GA	Gabon	MR	Mauritania
BE	Belgium	GB	United Kingdom	MW	Malawi
BF	Burkina Faso	GN	Guinea	NL	Netherlands
BG	Bulgaria	GR	Greece	NO	Norway
BJ	Benin	HU	Hungary	PL	Poland
BR	Brazil	IE	Ireland	RO	Romania
CA	Canada	IT	Italy	RU	Russian Federation
CF	Central African Republic	JP	Japan	SD	Sudan
CG	Congo	KP	Democratic People's Republic of Korea	SE	Sweden
CH	Switzerland	KR	Republic of Korea	SN	Senegal
CI	Côte d'Ivoire	LJ	Liechtenstein	SU	Soviet Union
CM	Cameroun	LK	Sri Lanka	TD	Chad
CS	Czechoslovakia	LU	Luxembourg	TC	Togo
DE	Germany	MC	Monaco	US	United States of America
DK	Denmark	MG	Madagascar		
ES	Spain				

REVISED
VERSION

INTERNATIONAL SEARCH REPORT

International Application No. PCT/US91/08606

I. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all) ³		
According to International Patent Classification (IPC) or to both National Classification and IPC		
IPC (5): A61K 37/36, 35/32; C07K 7/10, 13/00, 3/20, 3/28, 15/06; C12P 21/00 US CL : 514/12, 801, 886, 530/399; 435/69.1, 172.3, 424/85.8; 436/46, 512		
II. FIELDS SEARCHED		
Minimum Documentation Searched ⁴		
Classification System	Classification Symbols	
U.S.	530/324, 387, 399, 412, 416, 808; 435/7.1, 69.1, 172.3, 240.2; 514/12, 801, 886; 424/85.8; 436/64, 512	
Documentation Searched other than Minimum Documentation to the extent that such Documents are included in the Fields Searched ⁵		
Cas On-Line and APS		
III. DOCUMENTS CONSIDERED TO BE RELEVANT¹⁴		
Category ⁶	Citation of Document ¹⁵ with indication, where appropriate, of the relevant passages ¹⁷	Relevant to Claim No. ¹⁸
Y	US. A. 4,816,561 (Todaro) 28 March 1989 see the entire document particularly the abstract and column 15, line 16 to column 16, line 41.	1-4, 7, 8, 28-35, 42, 47-50
Y	US. A. 4,806,523 (Bentz et al) 21 February 1989, see the entire document particularly column 7, line 43 to column 8, line 13.	1,9-26
Y	US. A. 4,886,747 (Derynck et al) 12 December 1989, see the entire document particularly abstract.	36-42
Y	WO. A 88/05788 (Lucas et al.) 11 August 1988, see the document particularly paragraph bridging pages 10 and 11.	1-26, 33, 34
Y	Proc. Natl. Acad. Sci., Vol. 85, issued July 1988, P.T. Dijke et al., "Identification of another member of the transforming growth factor type B gene family", page 4715-4719, see the entire document, particularly the abstract on page 4715.	1-4, 36-41
<p>¹⁶ Special categories of cited documents: "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier document but published on or after the international filing date "L" document which may throw doubts on priority claims; or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed</p>		<p>¹⁷ later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "Z" document member of the same patent family</p>
IV. CERTIFICATION		
Date of the Actual Completion of the International Search ² 27 MAY 1992	Date of Mailing of this International Search Report ² 26 JUN 1992	
International Searching Authority ¹ ISA/US	Signature of Authorized Officer ²⁰ FATEMEH T. MOEZIE  LG	

I FURTHER INFORMATION CONTINUED FROM THE SECOND SHEET

Y

Science, Volume 249, issued 06 July 1990, Lefert et al., "Mediation of Cardioprotection by Transforming Growth-Factor-B", pages 61-64. See the entire document particularly abstract on page 61.

46

V. OBSERVATIONS WHERE CERTAIN CLAIMS WERE FOUND UNSEARCHABLE¹

1. Claim numbers , because they relate to subject matter (1) not required to be searched by this Authority, namely:

2. Claim numbers , because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out (1), specifically:

3. Claim numbers , because they are dependent claims not drafted in accordance with the second and third sentences of PCT Rule 8.4(a).

VI. OBSERVATIONS WHERE UNITY OF INVENTION IS LACKING²

This International Searching Authority found multiple inventions in this international application as follows:
SEE ATTACHED SHEET.

1. As all required additional search fees were timely paid by the applicant, this International search report covers all searchable claims of the international application.
2. As only some of the required additional search fees were timely paid by the applicant, this International search report covers only those claims of the international application for which fees were paid, specifically claims:
3. No required additional search fees were timely paid by the applicant. Consequently, this International search report is restricted to the invention first mentioned in the claims; it is covered by claim numbers:
4. As all searchable claims could be searched without effort justifying an additional fee, the International Search Authority did not invite payment of any additional fee.

Remark on protest:

- The additional search fees were accompanied by applicant's protest.
- No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

International Application No. PCT/US91/08606

I. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all)³

According to International Patent Classification (IPC) or to both National Classification and IPC

IPC (5): A61K 37/36; C07K 7/10, 13/00, 3/20
US CL : 530/324, 399, 412, 416; 514/021, 012

II. FIELDS SEARCHED

Minimum Documentation Searched⁴

Classification System	Classification Symbols
U.S.	530/324, 399, 412, 416; 514/021, 012

Documentation Searched other than Minimum Documentation⁵
to the extent that such Documents are included in the Fields Searched⁶

CAS ONLINE and APS

III. DOCUMENTS CONSIDERED TO BE RELEVANT¹⁴

Category ¹⁵	Citation of Document, ¹⁶ with indication, where appropriate, of the relevant passages ¹⁷	Relevant to Claim No. ¹⁸
Y	WO, A, 88/05788, LUCAS, et al. 11 August 1988, see the entire document, particularly paragraph bridging pages 10 and 11 and the paragraph on page 11.	1-7
Y	Proc. Natl. Acad. Sci., Vol. 85, issued July 1988, P.T. DUKE et al., "Identification of another member of the transforming growth factor type B gene family", pages 4715-4719, see the entire document, particularly the abstract on page 4715.	1-4
A	Molecular Endocrinology, Vol. 2, No. 8, issued 1988, "Complementary Deoxyribonucleic Acid Cloning of a Novel Transforming Growth Factor-B Messenger Ribonucleic Acid from Chick Embryo Chondrocytes", pages 747-755, see page 753, first column, first paragraph.	1-4

* Special categories of cited documents:¹⁹

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

- "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step
- "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
- "&" document member of the same patent family

IV. CERTIFICATION

Date of the Actual Completion of the International Search²

22 FEBRUARY 1992

Date of Mailing of this International Search Report²

17 MARCH 1992

International Searching Authority¹

ISA/US

Signature of Authorized Officer²⁰

Fatemeh Moezie

FURTHER INFORMATION CONTINUED FROM THE SECOND SHEET

Attachment to PCT/ISA/210
Invitation

Restriction of the following inventions is required

I. Claims 1-4 and 6, drawn to TGF-beta 2.3 polypeptide and a therapeutic composition, classified in Class 530, subclass 399.

Claim 5, drawn to a first method of preparing TGF-BETA 2.3 from bone, classified in Class 514, subclass 412.

Claim 7, drawn to a first method of inducing cartilage and/or bone formation, classified in Class 514, subclass 520, for example.

II. Claims 36-41, drawn to a method of preparing TGF-beta 2.3 recombinantly, classified in Class 435, subclass 172.3 for example.

III. Claim 8, drawn to a method of treating osteoporosis, classified in Class 514, subclass 12, for example.

IV . Claims 9-26, drawn to a method of treating, preventing or reducing inflammation, classified in Class 514, subclass 12, for example.

V . Claims 27, 43-46, drawn to a method of treating malfunction of hematopoiesis, classified in Class 514, subclass 577, for example.

VI . Claims 28-32, drawn to a method of treating the growth of a tumor cell, classified in Class 514, subclass 12, for example.

**This Page is Inserted by IFW Indexing and Scanning
Operations and is not part of the Official Record**

BEST AVAILABLE IMAGES

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images include but are not limited to the items checked:

- BLACK BORDERS**
- IMAGE CUT OFF AT TOP, BOTTOM OR SIDES**
- FADED TEXT OR DRAWING**
- BLURRED OR ILLEGIBLE TEXT OR DRAWING**
- SKEWED/SLANTED IMAGES**
- COLOR OR BLACK AND WHITE PHOTOGRAPHS**
- GRAY SCALE DOCUMENTS**
- LINES OR MARKS ON ORIGINAL DOCUMENT**
- REFERENCE(S) OR EXHIBIT(S) SUBMITTED ARE POOR QUALITY**
- OTHER:** _____

IMAGES ARE BEST AVAILABLE COPY.

As rescanning these documents will not correct the image problems checked, please do not report these problems to the IFW Image Problem Mailbox.